TABLE 7

	Summary of change from Baseline in IBS QOL (32 µg)									
	Component/Scale Score									
	QOL Overall	Dysphoria	Interference with Activity	Body Image	Health Worry	Food Avoidance	Social Reaction	Sexual	Relationship	
Baseline	_									
N Mean (Std) Week 4	49 60.14 22.05	49 59.69 24.79	49 69.82 23.385	49 43.37 24.387	49 44.38 24.672	49 52.21 32.175	49 66.84 28.562	49 68.62 31.367	49 70.23 23.385	
N Mean (Std) Week 12	36 11.25** 16.277	36 14.58** 20.39	36 7.04* 17.753	36 15.28** 18.264	36 16.9** 17.534	36 8.33* 23.988	36 9.9** 16.46	36 8.68** 18.131	36 7.64* 18.831	
N Mean (Std) End of Study	33 13.08** 13.527	33 15.25** 15.285	33 9.42** 18.052	33 17.99** 15.21	33 19.44** 21.616	33 11.36** 23.46	33 10.8** 14.173	33 10.98** 18.424	33 9.09* 20.87	
N Mean (Std)	44 12.58** 12.621	44 14.77** 14.429	44 8.2** 16.726	44 17.47** 15.344	44 17.61** 22.317	44 10.79** 21.906	44 11.08** 14.32	44 10.23** 16.894	44 10.79** 20.139	

TABLE 8

	Summary of change from Baseline in IBS QOL (48 µg)									
	Component/Scaled Score									
	QOL Overall	Dysphoria	Interference with Activity	Body Image	Health Worry	Food Avoidance	Social Reaction	Sexual	Relationship	
Baseline	_									
N Mean (Std) Week 4	45 59.85 21.664	45 56.81 26.802	45 68.25 23.396	45 45.69 21.396	45 44.07 23.274	45 50.37 31.927	45 66.81 27.074	45 71.94 30.404	45 75.18 22.365	
N Mean (Std) Week 12	34 12.43** 11.619	34 17.28** 16.842	34 9.14** 14.477	34 13.24** 13.568	34 19.61** 18.562	34 12.01** 19.59	34 8.82** 14.028	34 8.46** 15.607	34 6.87** 12.558	
N Mean (Std) End of Study	30 14.8** 13.65	30 20.83** 18.863	30 10.95** 15.091	30 17.08** 18.419	30 23.33** 18.098	30 11.3**9 22.153	30 14.17** 21.143	30 5 18.159	30 6.95** 12.202	
N Mean (Std)	43 11.54** 13.002	43 16.28** 18.62	43 7.97** 14.387	43 14.24** 17.43	43 17.05** 19.067	43 8.33** 20.002	43 12.06** 18.72	43 3.49 17.535	43 6.01** 12.244	

<sup>\*</sup>P < 0.05,

The description of the invention is merely exemplary in  $_{60}$  nature and, thus, variations that do not depart from the gist of the invention are intended to be within the scope of the invention. Such variations are not to be regarded as a departure from the spirit and scope of the invention.

All patents and publications cited in this specification are herein incorporated by reference

What is claimed is:

1. A method for the long term treatment of chronic constipation in a human subject, wherein the treatment comprises administering to the subject in need thereof an effective amount of a prostaglandin compound represented by Formula (I) and/or its tautomer:

<sup>\*\*</sup>P < 0.01 (paired T-Tests)